

CLAIMS

What is claimed:

1. (Previously amended) A method of treating a subject suffering from anemia comprising administering a therapeutically effective amount of a human antibody, or an antigen-binding fragment thereof, to the subject, wherein the antibody dissociates from human TNF α with a K_d of 1×10^{-8} M or less and a K_{off} rate constant of 1×10^{-3} s $^{-1}$ or less, both determined by surface plasmon resonance, and neutralizes human TNF α cytotoxicity in a standard *in vitro* L929 assay with an IC $_{50}$ of 1×10^{-7} M or less, such that the anemia is treated.
2. (Previously amended) A method of treating a subject suffering from anemia comprising administering a therapeutically effective amount of a human antibody, or an antigen-binding fragment thereof, with the following characteristics:
 - a) dissociates from human TNF α with a K_{off} rate constant of 1×10^{-3} s $^{-1}$ or less, as determined by surface plasmon resonance;
 - b) has a light chain CDR3 domain comprising the amino acid sequence of SEQ ID NO: 3, or modified from SEQ ID NO: 3 by a single alanine substitution at position 1, 4, 5, 7 or 8 or by one to five conservative amino acid substitutions at positions 1, 3, 4, 6, 7, 8 and/or 9;
 - c) has a heavy chain CDR3 domain comprising the amino acid sequence of SEQ ID NO: 4, or modified from SEQ ID NO: 4 by a single alanine substitution at position 2, 3, 4, 5, 6, 8, 9, 10 or 11 or by one to five conservative amino acid substitutions at positions 2, 3, 4, 5, 6, 8, 9, 10, 11 and/or 12.
3. (Previously amended) A method of treating a subject suffering from anemia comprising administering a therapeutically effective amount of a human antibody, or an antigen-binding fragment thereof, with a light chain variable region (LCVR) comprising the amino acid sequence of SEQ ID NO:1 and a heavy chain variable region (HCVR) comprising the amino acid sequence of SEQ ID NO:2.
4. (Currently amended) The method of any one of claims 1, 2, ~~and or~~ 3, wherein the antibody, or ~~antigen-binding fragment thereof~~, is D2E7.

5. (Currently amended) The method of any one of claims 1, 2, ~~and~~ or 3, wherein anemia is selected from the group consisting of: anemia related to rheumatoid arthritis, anemia of infection and chronic inflammatory diseases, iron deficiency anemia, autoimmune hemolytic anemia, myelophthisic anemia, aplastic anemia, hypoplastic anemia, pure red cell aplasia, anemia associated with renal failure or endocrine disorders, megaloblastic anemias, defects in heme or globin synthesis, sickle-cell anemia, sideroblastic anemia, anemia associated with chronic infections, and myelophthisic anemias.

6. (Previously amended) A method for inhibiting human TNF α activity in a human subject suffering from anemia comprising administering a therapeutically effective amount of a human antibody, or an antigen-binding fragment thereof, to the subject, wherein the antibody dissociates from human TNF α with a K_d of 1×10^{-8} M or less and a K_{off} rate constant of 1×10^{-3} s $^{-1}$ or less, both determined by surface plasmon resonance, and neutralizes human TNF α cytotoxicity in a standard *in vitro* L929 assay with an IC_{50} of 1×10^{-7} M or less.

7. (Original) The method of claim 6, wherein anemia is selected from the group consisting of: anemia related to rheumatoid arthritis, anemia of infection and chronic inflammatory diseases, iron deficiency anemia, autoimmune hemolytic anemia, myelophthisic anemia, aplastic anemia, hypoplastic anemia, pure red cell aplasia, anemia associated with renal failure or endocrine disorders, megaloblastic anemias, defects in heme or globin synthesis, sickle-cell anemia, sideroblastic anemia, anemia associated with chronic infections, and myelophthisic anemias.

8. (Currently amended) The method of any one of claims 6 ~~and~~ or 7, wherein the antibody, ~~or antigen binding fragment thereof~~, is D2E7.

9. (Original) A method of treating a subject suffering from anemia comprising administering a therapeutically effective amount of D2E7, or an antigen-binding fragment thereof, to the subject, such that anemia is treated.

10. (Original) The method of claim 9, wherein anemia is selected from the group consisting of: anemia related to rheumatoid arthritis, anemia of infection and chronic inflammatory diseases, iron deficiency anemia, autoimmune hemolytic anemia, myelophthisic anemia, aplastic anemia, hypoplastic anemia, pure red cell aplasia, anemia associated with renal failure or endocrine disorders, megaloblastic

anemias, defects in heme or globin synthesis, sickle-cell anemia, sideroblastic anemia, anemia associated with chronic infections, and myelophthisic anemias.

11. (Original) A method of treating a subject suffering from anemia comprising administering a therapeutically effective amount of D2E7, or an antigen-binding fragment thereof, and at least one additional therapeutic agent to the subject, such that anemia is treated.

12 (Withdrawn) A kit comprising:

a) a pharmaceutical composition comprising a human antibody, or an antigen binding portion thereof, and a pharmaceutically acceptable carrier, wherein the antibody dissociates from human TNF α with a K_d of 1×10^{-8} M or less and a K_{off} rate constant of 1×10^{-3} s $^{-1}$ or less, both determined by surface plasmon resonance, and neutralizes human TNF α cytotoxicity in a standard *in vitro* L929 assay with an IC_{50} of 1×10^{-7} M or less; and

b) instructions for administering to a subject the TNF α antibody pharmaceutical composition for treating a subject who is suffering from anemia.

13. (Withdrawn) A kit according to claim 12, wherein the antibody, or an antigen binding portion thereof, is D2E7.